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SECTION I - REQUEST FOR APPROVAL OF THE FOLLOWING ITEMS (This section will be initiated by the contractor)								
TO: Eastern Environmental Project Office USAED Seattle 1910 Northwest Boulevard, Suite 210 Coeur d'Alene, ID 83814			FROM: Amec Foster Wheeler Environment & Infra 751 Arbor Way, Ste 180 Blue Bell, PA 19422-1972		CONTRACT NO. <div style="text-align: center;">W912DW-16-C-0012 NA</div>		CHECK ONE: <input checked="" type="checkbox"/> THIS IS A NEW TRANSMITTAL <input type="checkbox"/> THIS IS A RESUBMITTAL OF TRANSMITTAL _____	
SPECIFICATION SEC. NO. (Cover only one section with each transmittal) 01 35 45.00 10 Chemical Data Quality Control			PROJECT TITLE AND LOCATION 01-Main Register CTP Upgrade and Ground Water Collection Bunker Hill Superfund Site, Kellogg, ID			THIS TRANSMITTAL IS FOR: (Check one) <input checked="" type="checkbox"/> MFG <input type="checkbox"/> ID <input type="checkbox"/> GA <input type="checkbox"/> DA <input type="checkbox"/> CR <input type="checkbox"/> DA/CR <input type="checkbox"/> DA/GA		
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e.	f.							
4.1	DMR June 2017	SD-06	0	3.7		A		
REMARKS				I certify that the above submitted items have been reviewed in detail and are correct and in the strict conformance with the contract drawings and specifications except as otherwise stated. <div style="text-align: right; margin-right: 50px;">  NAME AND SIGNATURE OF CONTRACTOR </div>				
SECTION II - APPROVAL ACTION								
ENCLOSURES RETURNED (List by item No.)			NAME, TITLE AND SIGNATURE OF APPROVING AUTHORITY				DATE	



AMEC FOSTER WHEELER
BUNKER HILL CENTRAL TREATMENT PLANT UPGRADE PROJECT
Kellogg, Idaho

TRANSMITTAL FORM: DMR - June 2017 Submittal

☒ Approved

☐ Approved with corrections as noted on submittal data and/or attached sheet(s)

Signature: 

By: Spencer Archer, PE

Title: Commissioning and O&M Manager

Date: July 14, 2017

☒ Approved

☐ Approved with corrections as noted on submittal data and/or attached sheet(s)

Signature: 

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Date: July 14, 2017

Distribution: USACE, Seattle District
Amec Foster Wheeler



DATA VALIDATION REPORT

Bunker Hill Central Treatment Plant

Kellogg, Idaho

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July 2017

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ACRONYMS

%	percent
°C	degrees Celsius
Amec Foster Wheeler	Amec Foster Wheeler Environment & Infrastructure, Inc.
CLP	Contract Laboratory Program
COC	chain of custody
CCV	continuing calibration verification
EPA	United States Environmental Protection Agency
ICAL	initial calibration
ICV	initial calibration verification
ID	identification
LCS	laboratory control sample
LCSD	laboratory control sample duplicate
LOQ	limit of quantitation
MDL	method detection limit
mg/L	milligrams per liter
MS	matrix spike
MSD	matrix spike duplicate
QC	quality control
RL	reporting limit
RPD	relative percent difference
SAP	sampling and analysis plan
SM	Standard Methods for the Examination of Water and Wastewater
SVL	SVL Analytical Inc.
TSS	total suspended solids

DATA VALIDATION REPORT

Bunker Hill Central Treatment Plant

Kellogg, Idaho

1.0 INTRODUCTION

Amec Foster Wheeler Environment & Infrastructure, Inc. (Amec Foster Wheeler) collected 26 water samples (including two field duplicates, one equipment blank, and one trip blank) between May 22 and June 26, 2017 from the Bunker Hill Central Treatment Plant in Kellogg, Idaho. Amec Foster Wheeler submitted the samples to SVL Analytical Inc. (SVL), located in Coeur D'Alene, Idaho, where they were assigned to sample delivery groups X7E0428, X7E0580, X7F0028, X7F0060, X7F0061, X7F0118, X7F0204, X7F0235, X7F0236, X7F0386, X7F0429, X7F0477, X7F0558, X7F0586, and X7F0645. SVL analyzed the samples for total metals by United States Environmental Protection Agency (EPA) Method 200.7, total suspended solids (TSS) by Standard Methods for the Examination of Water and Wastewater (SM) 2540D; and pH by SM 4500B. A list of these samples by field sample identification (ID), sample collection date, and the laboratory sample IDs is presented in Table 1.

2.0 DATA VALIDATION METHODOLOGY

Amec Foster Wheeler performed Stage 4 validation on samples KT-05-22-17, KT-05-25-17, and PTM-05-25-17. The Stage 4 validation includes review and recalculation of the laboratory's analytical report and the raw analytical data. The remainder of the data underwent EPA Stage 2B validation, which includes review of sample- and instrument-specific quality control (QC) samples on data summary forms, but does not include review or validation of the raw analytical data. This data validation has been performed in general accordance with:

- Amec Foster Wheeler, 2017. Operations & Maintenance Services Sampling and Analysis Plan (SAP), Bunker Hill Central Treatment Plant Upgrade Project, Kellogg, Idaho, March 2017.
- EPA, 2014. EPA Contract Laboratory Program (CLP) National Functional Guidelines for Inorganic Superfund Data Review, EPA-540 R 013 001.
- The analytical methods referenced by the laboratory.

The laboratory's certified analytical report and supporting documentation were reviewed to assess the following:

- Data package and electronic data deliverable completeness;
- Chain-of-custody (COC) compliance;
- Sample Receipt
- Holding time compliance;
- Initial calibration (ICAL), initial calibration verification (ICV), and continuing calibration verification (CCV) compliance with method specified criteria;
- Presence or absence of laboratory contamination as demonstrated by calibration and laboratory blanks;
- Accuracy and bias as demonstrated by recovery of surrogate spikes, laboratory control sample (LCS), and matrix spike (MS) samples;
- Analytical precision as relative percent difference (RPD) of analyte concentration between laboratory duplicates, LCSs/LCS duplicates (LCSs), or MS/MS duplicates (MSDs);
- Sampling and analytical precision as RPD of analyte concentration between field duplicates;
- Internal standard and surrogate compound recoveries;
- Analyte identification and quantification verification from raw analytical data (Stage 4 validation only); and
- Insofar as possible, the degree of conformance to method requirements and good laboratory practices

3.0 EXPLANATION OF DATA QUALITY INDICATORS

Summary explanations of the specific data quality indicators reviewed during data validation are presented below.

3.1 LABORATORY CONTROL SAMPLE RECOVERIES

LCSs are aliquots of analyte free matrices that are spiked with the analytes of interest for an analytical method, or a representative subset of those analytes. The spiked matrix is then processed through the same analytical procedures as the samples they accompany. LCS recovery is an indication of a laboratory's ability to successfully perform an analytical method in an interference free matrix.

3.2 MATRIX SPIKE RECOVERIES

MSs and MSDs are prepared by adding known amounts of the analytes of interest for an analytical method, or a representative subset of those analytes, to an aliquot of sample. The spiked sample is then processed through the same extraction, concentration, cleanup, and analytical procedures as the unspiked samples in an analytical batch.

MS recovery and precision are an indication of a laboratory's ability to successfully recover an analyte in the matrix of a specific sample or closely related sample matrices. It is important not to apply MS results for any specific sample to other samples without understanding how the sample matrices are related.

3.3 BLANK CONCENTRATIONS

Blank samples are aliquots of analyte free matrix that are used as negative controls to verify that the sample collection, storage, preparation, and analysis system does not produce false positive results.

Laboratory blanks are processed by the laboratory using exactly the same procedures as the field samples. Target analytes should not be found in laboratory blanks.

Equipment blanks are prepared by passing analyte-free water through or over sample collection equipment and collecting the water in sample containers. Equipment blanks are analyzed for the analytical suite required for the project. Equipment blanks are used to monitor for possible sample contamination during the sample collection process and serve as a check on the effectiveness of field decontamination procedures.

Trip blanks are vials of analyte free water that accompany sample bottles shipped to the field and back to the laboratory with field samples. Trip blanks assess contamination attributed to shipping and handling procedures, as well as contamination from containers. Target analytes should not be found in trip blanks.

Target analytes should not be found in laboratory, equipment, and trip blanks. When target analytes are detected in blanks, analyte concentrations in associated samples less than five times the concentration detected in the blank will be U qualified as being not detected.

3.4 LABORATORY DUPLICATES

Laboratory duplicate analysis verifies acceptable method precision by the laboratory at the time of preparation and analysis and/or sampling precision at the time of collection.

4.0 DEFINITIONS OF QUALIFIERS THAT MAY BE ADDED DURING DATA VALIDATION

- J** The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- R** The sample result is rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.
- U** The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- UJ** The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.

5.0 QUALIFICATION REASON CODES

The following reason codes were applied to the data during data validation:

- DL** The analyte concentration is between the method detection limit (MDL) and the reporting limit (RL).
- HT** Holding time exceeded.

6.0 CHAIN OF CUSTODY AND SAMPLE RECEIPT CONDITION DOCUMENTATION

The samples were received at the laboratory under proper COC, intact, properly preserved, and at temperatures less than the SAP-specified maximum of 6 degrees Celsius (°C), with the following exception:

- According to SVL's sample receipt documentation, sample 006-06-26-17 was received at a temperature of 6.3 C. The laboratory received the sample on ice within eight hours of sampling and data usability is not adversely affected.

7.0 SPECIFIC DATA VALIDATION FINDINGS

Results from these samples may be considered usable with the limitations and exceptions described in Sections 7.1 through 8.0. Qualifiers added during validation are summarized in Table 2.

7.1 METALS BY EPA METHOD 200.7

Total metals results generated by SVL may be considered usable with the limitations described in Sections 7.1.1 through 7.1.9.

7.1.1 Holding Times

All samples were analyzed for metals within the SAP-specified holding of 180 days

7.1.2 Initial and Continuing Calibration Verification

ICV and CCV recoveries were within method-specified limits.

7.1.3 Initial and Continuing Calibration Blanks

Target analytes were not detected in the initial calibration blanks and continuing calibration blanks.

7.1.4 Laboratory, Equipment, and Trip Blanks

Target analytes were not detected in the laboratory, equipment, and trip blanks associated with the analysis of these samples, with the following exceptions:

- Zinc was detected at concentrations of 0.038 milligrams per liter (mg/L) and 0.001 mg/L, respectively, in the trip blank and equipment blank associated with sample KT-06-05-17 and its field duplicate, QC-06-05-17. Zinc was detected in the associated samples at concentrations greater than 5 times the detections in the associated blanks, and data usability is not adversely affected.

7.1.5 Laboratory Control Sample Accuracy and Precision

LCS and LCSD recoveries were within the SAP-specified limits, and RPDs between the LCS and LCSD results were less than the SAP-specified maximum of 20 percent (%).

7.1.6 Laboratory Duplicates

SVL performed duplicate analyses on project samples PTM-05-25-17, 006-06-02-17, 006-06-05-17, 006-06-07-17, KT-05-29-17, 006-06-09-17, 006-06-12-17, TB-05-06-17, 006-06-14-17, 006-06-16-17, 006-06-19-17, 006-06-21-17, KT-06-12-17, 006-06-23-17, 006-06-26-17 and 006-06-28-17. RPDs between source and duplicate results were less than the SAP-specified maximum of 20%, with the following exception:

- The RPD for lead was high at 24.2% in the duplicate analysis of sample PTM-05-25-17. The difference between the primary and duplicate results was less than the RL, demonstrating acceptable analytical precision.

7.1.7 Matrix Spikes/Matrix Spike Duplicates

SVL performed MS and MSD analyses on project samples PTM-05-25-17, 006-06-02-17, 006-06-05-17, 006-06-07-17, KT-05-29-17, 006-06-09-17, 006-06-12-17, TB-06-05-17, 006-06-14-17, 006-06-16-17, 006-06-19-17, 006-06-21-17, KT-06-12-17, 006-06-23-17, 006-06-26-17, and 006-06-28-17 for total metals by EPA method 200.7. MS/MSD recoveries were within SAP-specified limits, and RPDs between MS and MSD results were less than the laboratory-specified maximum of 20%, with the following exceptions:

- Manganese (167%, MS) and zinc (373%, 205%) recoveries were high in the MS and/or MSD performed on sample KT-05-29-17. Manganese (99.0 mg/L) and zinc (184 mg/L) were detected in the unspiked native sample at concentrations greater than four times the spike concentration of 1.0 mg/L, and data usability cannot be evaluated based on the MS/MSD performance of these analytes in this sample.
- Manganese recovery was low at 50.9% in the MSD performed on sample 006-06-12-17. The manganese concentration detected in the unspiked native sample, at 6.29 mg/L, was more than four times the spike concentration of 1.0 mg/L and it is not possible to evaluate data usability for this analyte in this sample based on MSD recovery.
- Manganese recoveries were low at 61.9% and 77.9%, respectively, in the MS and MSD performed on sample 006-06-16-17. The manganese concentration detected in the unspiked native sample, at 6.32 mg/L, was more than four times the spike concentration of 1.0 mg/L and it is not possible to evaluate data usability for this analyte in this sample based on MS/MSD recovery.
- Manganese recovery was low at 43.6% in the MSD performed on sample 006-06-19-17. The manganese concentration detected in the unspiked native sample, at 6.06 mg/L, was more than four times the spike concentration of 1.0 mg/L and it is not possible to evaluate data usability for this analyte in this sample based on MSD recovery.

- Manganese recoveries were high at 123% and 123%, respectively, in the MS and MSD performed on sample 006-06-21-17. The manganese concentration detected in the unspiked native sample, at 5.09 mg/L, was more than four times the spike concentration of 1.0 mg/L and it is not possible to evaluate data usability for this analyte in this sample based on MSD recovery.
- Manganese (-60.1%, 31.6%) and zinc (-63.0%, -5.00%) recoveries were low in the MS and MSD performed on sample KT-06-12-17. Manganese (88.3 mg/L) and zinc (138 mg/L) were detected in the native unspiked sample at concentrations greater than four times the spike concentration of 1.0 mg/L, and data usability cannot be evaluated based on the MS/MSD performance of these analytes in this sample.
 - Manganese recovery was low at -63.4% in the MS performed on sample 006-06-23-17. The manganese concentration detected in the unspiked native sample, at 4.56 mg/L, was more than four times the spike concentration of 1.0 mg/L and it is not possible to evaluate data usability for this analyte in this sample based on MSD recovery.
 - Manganese recovery was high at 131% in the MS performed on sample 006-06-26-17. The manganese concentration detected in the unspiked native sample, at 8.30 mg/L, was more than four times the spike concentration of 1.0 mg/L and it is not possible to evaluate data usability for this analyte in this sample based on MS recovery.

7.1.8 Analytical Sensitivity

Amec Foster Wheeler compared RLs for cadmium, lead, manganese, and zinc against applicable discharge limits to confirm that the RLs were sufficiently low to meet the discharge limits. Non-detect results were reported to RLs less than the applicable discharge limits.

7.1.9 Data Reporting and Analytical Procedures

SVL J qualified analytes with concentrations between the MDL and the RL. Amec Foster Wheeler agrees that these results are quantitatively uncertain and has maintained SVL's J qualifiers. (J-DL)

7.2 TOTAL SUSPENDED SOLIDS BY SM 2540D

TSS results generated by SVL may be considered fully usable without qualification.

7.2.1 Holding Times

All samples were analyzed for TSS within the SAP-specified holding time of 7 days.

7.2.2 Laboratory Blanks

TSS was not detected in the laboratory blanks associated with the analysis of these samples.

7.2.3 Laboratory Control Sample Accuracy and Precision

LCS and LCSD recoveries were within the laboratory-specified 90 to 110% limits and RPDs between the LCS and LCSD results were less than the laboratory-specified maximum of 10%.

7.2.4 Laboratory Duplicates

SVL performed duplicate analyses on project samples 006-06-02-17, 006-06-05-17, 006-06-07-17, KT-05-29-17, 006-06-09-17, 006-06-12-17, PTM-06-08-17, 006-06-14-17, 006-06-16-17, 006-06-19-17, 006-06-21-17, KT-06-12-17, 006-06-23-17, 006-06-26-17, and 006-06-28-17. RPDs between source and duplicate results were less than laboratory-specified maximum of 10%, with the following exceptions:

- RPDs for TSS were high in the duplicate analyses of samples PTM-06-08-17 (40.0%), 006-06-14-17 (35.3%), 006-06-16-17 (25.0%), and 006-06-28-17 (82.35%). The differences between primary and duplicate results were less than the RL, demonstrating acceptable analytical precision.

7.2.5 Analytical sensitivity

Amec Foster Wheeler compared RLs for TSS against applicable discharge limits to confirm that the RLs were sufficiently low to meet the discharge limits. Non-detect results were reported to RLs less than the applicable discharge limits.

7.2.6 Data Reporting and Analytical Procedures

There were no data anomalies associated with the TSS analysis of these samples.

7.3 pH BY SM 4500B

pH results generated by SVL may be considered usable with the limitations described in Sections 7.3.1 through 7.3.4.

7.3.1 Holding Times

All samples were analyzed for pH after the method-specified 15-minute hold time had passed. Amec Foster Wheeler J qualified the pH results from these samples because of the missed hold time. (J-HT)

7.3.2 Laboratory Control Sample Accuracy

LCS recoveries were within the laboratory-specified 98.5 to 101.5% limits.

7.3.3 Laboratory Duplicates

SVL performed duplicate analyses on samples 006-06-02-17, 006-06-05-17, 006-06-07-17, 006-06-09-17, 006-06-12-17, 006-06-14-17, 006-06-16-17, 006-06-19-17, 006-06-21-17, KT-06-15-17, 006-06-23-17, 006-06-26-17 and 006-06-28-17. RPDs between source and duplicate results were less than laboratory-specified maximum of 5%.

7.3.4 Data Reporting and Analytical Procedures

There were no data anomalies associated with the pH analysis of these samples.

8.0 FIELD DUPLICATES

Field duplicates were collected with samples KT-06-05-17 (QC-06-05-17) and 006-06-28-17 (QC-06-28-17). Target analyte detections are summarized in Table 3. Precision values were less than the QAPP-specified maximum of 30%, or the differences between detected concentrations were less than the limit of quantitation (LOQ), demonstrating acceptable sampling and analytical precision.

9.0 SUMMARY AND CONCLUSIONS

Amec Foster Wheeler reviewed 132 data records from field samples during this validation. All the data generated are usable and of acceptable quality with the addition of qualifiers presented in Table 2. Qualifier definitions are summarized in Section 4.0, reason codes are summarized in Section 5.0, and qualified data are summarized below.

- Amec Foster Wheeler J qualified 15 records (11.3%) as being estimated concentrations because of hold time exceedances or analyte concentrations between the MDL and RL.

No records were rejected and 100% of the data should be considered valid with the addition of the qualifiers presented in Table 2.

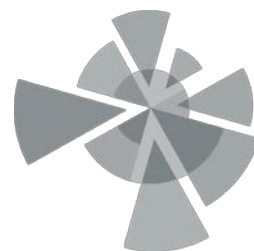
REFERENCES

Amec Foster Wheeler, 2017. Operations & Maintenance Services Sampling and Analysis Plan (SAP), Bunker Hill Central Treatment Plant Upgrade Project, Kellogg, Idaho, March 2017.

EPA, 2014. EPA Contract Laboratory Program (CLP) National Functional Guidelines for Inorganic Superfund Data Review, EPA-540 R 013 001.

LIMITATIONS

This report was prepared exclusively for the Bunker Hill Central Treatment Plant by Amec Foster Wheeler Environment & Infrastructure, Inc. The quality of information, conclusions, and estimates contained herein is consistent with the level of effort involved in Amec Foster Wheeler services and based on: i) information available at the time of preparation, ii) data supplied by outside sources, and iii) the assumptions, conditions, and qualifications set forth in this report. This data validation report is intended to be used by the Bunker Hill Central Treatment Plant in Kellogg, Idaho only, subject to the terms and conditions of its contract with Amec Foster Wheeler. Any other use of, or reliance on, this report by any third party is at that party's sole risk.



TABLES

TABLE 1
Field Samples Submitted to SVL Analytical, Inc.
Bunker Hill Central Treatment Plant Upgrade Program
Kellogg, Idaho

Field Sample ID	Collection Date	SVL Analytical, Inc. Sample ID	Notes
KT-05-22-17	5/22/2017	X7E0428-01	Stage 4 Validation
KT-05-25-17	5/25/2017	X7E0428-02	Stage 4 Validation
PTM-05-25-17	5/25/2017	X7E0428-03	Stage 4 Validation
KT-05-29-17	5/29/2017	X7E0580-01	
KT-06-01-17	6/1/2017	X7E0580-02	
006-06-02-17	6/2/2017	X7F0028-01	
006-06-05-17	6/5/2017	X7F0060-01	
KT-06-05-17	6/5/2017	X7F0061-01	
QC-06-05-17	6/5/2017	X7F0061-02	Field Duplicate of KT-06-05-17
TB-06-05-17	6/5/2017	X7F0061-03	Trip Blank
RB-06-05-17	6/5/2017	X7F0061-04	Equipment Blank
KT-06-08-17	6/8/2017	X7F0061-05	
PTM-06-08-17	6/8/2017	X7F0061-06	
006-06-07-17	6/7/2017	X7F0118-01	
006-06-09-17	6/9/2017	X7F0204-01	
006-06-12-17	6/12/2017	X7F0235-01	
KT-06-12-17	6/12/2017	X7F0236-01	
KT-06-15-17	6/15/2017	X7F0236-02	
006-06-14-17	6/14/2017	X7F0308-01	
006-06-16-17	6/16/2017	X7F0386-01	
006-06-19-17	6/19/2017	X7F0429-01	
006-06-21-17	6/21/2017	X7F0477-01	
006-06-23-17	6/23/2017	X7F0558-01	
006-06-26-17	6/26/2017	X7F0586-01	
006-06-28-17	6/28/2017	X7F0645-01	
QC-06-28-17	6/28/2017	X7F0645-02	Field Duplicate of 006-06-28-17

Notes:

ID = identification

TABLE 2
Qualifiers Added During Data Validation
Bunker Hill Central Treatment Plant Upgrade Program
Kellogg, Idaho

Sample IDs	Methods	Analytes	Concentrations	Qualifiers and Reason Codes	
006-06-02-17	4500H	pH	7.40 SU	J	HT
006-06-05-17	4500H	pH	7.19 SU	J	HT
006-06-07-17	EPA 200.7	Lead	0.0026 mg/L	J	DL
	4500H	pH	7.13 SU	J	HT
006-06-14-17	4500H	pH	7.11 SU	J	HT
006-06-16-17	4500H	pH	7.03 SU	J	HT
006-06-19-17	4500H	pH	7.01 SU	J	HT
006-06-21-17	4500H	pH	7.11 SU	J	HT
006-06-23-17	4500H	pH	7.17 SU	J	HT
006-06-26-17	4500H	pH	7.17 SU	J	HT
KT-06-15-17	4500H	pH	2.84 SU	J	HT
PTM-06-08-17	EPA 200.7	Lead	0.0050 mg/L	J	DL
RB-06-05-17	EPA 200.7	Zinc	0.001 mg/L	J	DL
006-06-28-17	4500H	pH	7.22 SU	J	HT
QC-06-28-17	4500H	pH	7.24 SU	J	HT

Notes:

SU = Standard Units

mg/L = milligrams per liter

Qualifier Definitions:

J = The analyte was positively identified; the associated numerical value is approximate.

UJ = The analyte was not detected above the RL. However, the RL is approximate.

Reason Codes:

DL = The analyte concentration is between the method detection limit (MDL) and the reporting limit (RL).

HT = Holding time exceedence

TABLE 3
Field Duplicate Detections
Bunker Hill Central Treatment Plant Upgrade Program
Kellogg, Idaho

Analyte	Method	Average LOQ	Primary Concentration	Field Duplicate	RPD	Notes
Samples KT-06-05-17 and QC-06-05-17						
Cadmium	EPA 200.7	0.002 mg/L	0.391	0.388	1%	
Lead	EPA 200.7	0.0075 mg/L	0.711	0.707	1%	
Manganese	EPA 200.7	0.008 mg/L	102	99.6	2%	
Zinc	EPA 200.7	0.1 mg/L	162	158	3%	
TSS	SM 2540 D	5 mg/L	99.0	97.0	2%	
Samples KT-06-05-17 and QC-06-05-17						
Cadmium	EPA 200.7	0.002 mg/L	0.005	0.005	0%	
Lead	EPA 200.7	0.0075 mg/L	0.008 U	0.008 U	NC	± LOQ
Manganese	EPA 200.7	0.008 mg/L	8.6	8.8	2%	
Zinc	EPA 200.7	0.1 mg/L	0.182	0.189	4%	
TSS	SM 2540 D	5 mg/L	1.0	1.0	0%	

Notes:

TSS = Total Suspended Solids

Qualifier and Reason Code Definitions:

U = The analyte was analyzed for, but was not detected above the reported sample quantitation limit.

± LOQ = The difference between analyte concentrations is less than the limit of quantification, demonstrating acceptable sampling and/or analytical precision.